# **Endocrine Disruptors Research Program Annual Performance Goals (APGs)**

APGs represent a program's major milestones toward accomplishing its long-term goals. The following charts outline the program's success in meeting its planned APGs on time.

### **FY 2003**

APG Title	<b>Completion Status</b>	Explanation
Develop tools to identify hazards and	Met as planned	N/A
evaluate existing approaches to manage		
risks from exposure to endocrine		
disrupting chemicals (EDCs) capable of		
inducing adverse effects in humans and		
wildlife.		

### FY 2004

112001		
APG Title	<b>Completion Status</b>	Explanation
Characterize sources of exposure and	Met as planned	N/A
environmental fates of EDCs.		
Determine efficacy of wildlife species	Met as planned*	N/A
as sentinels.	_	
Evaluate existing risk management tools to reduce exposure to EDCs.	Delayed by one fiscal year - the APG was subsequently completed in late 2005.	There were insufficient personnel to complete the supporting APM in FY 2004 because of personnel loss due to 1) retirement and 2) details to acting positions outside of the Division. The delay did not impact any regulatory actions.
Evaluate several classes of chemicals suspected of being EDCs and determine potencies in laboratory studies.	Met as planned	N/A
Evaluate several classes of chemicals suspected of being EDCs in field studies and ascertain degree to which they adversely affect wildlife at the population level.	Met as planned*	N/A

<sup>\*</sup> Note: Due to the closure of NHEERL's Gulf Breeze Division as a result of Hurricane Ivan, data on the completion of these APGs were not available at the end of the FY 2004. However, the program subsequently determined that the APGs were in fact completed within FY 2004, and marked the APGs complete in ORD's system as of March, 2005.

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## FY 2005

APG Title	Completion Status	s Explanation
Determine the shape of the dose-	Met as planned	N/A
response curve in a variety of species		
exposed to ambient levels of EDCs.		

## FY 2006

APG Title	Completion Status	Explanation
Determine critical biological factors	Met as planned	N/A
during development resulting in		
toxicities later in life.		
Determine degree to which effects of	Met as planned	N/A
EDCs with defined mechanisms/modes		
of action can be extrapolated across		
classes of vertebrates.		
Determine extent to which exposure to	Met as planned	N/A
EDCs contribute to onset or increase in		
severity of diseases.		
Develop standardized protocols for	Met as planned	N/A
screening chemicals for their potential		
endocrine-mediated effects to meet		
FQPA requirements.		